

# TRD Sr. QA Specialist

Job ID

REQ-10074537

Mar 23, 2026

LOC\_IT

## About the Role

Major Accountabilities:

- Support site qualification and validation activities (advising, review, approval).
- Implementation of Quality Systems (incl. documentation management)
- Supplier management activities (agreements, oversight, audit).
- Preparation/support and coordination of CAPA/follow -up
- Audit and inspection preparation and support, ensure applications, certificate maintenance etc. to local HA
- Change control review/approval
- Ensure local DI and eCompliance oversight (training, inspections, plan, risk ID etc)
- KPI/KQI trending
- Handling of technical complaints, deviations, quality events related to Novartis products, systems or processes.

Key Performance Indicators:

- Successful support of projects with agreed quality and delivery dates, passing of internal & external inspections.
- Meet quality & timelines for all projects
- Act in accordance with Novartis standards.
- The number and severity of cGMP issues identified during internal and external audits
- Year-end figures within budget; Successful coordination of departmental operational activities

Work Experience:

- Change Control Management
- Audit & Inspection Management
- Compliance Risk Management
- Good Manufacturing Practices (cGMP)
- GxP Experience
- KPI Reporting
- Quality Management System

Prerequisites:

- Minimum of 5 years in pharmaceutical industry (sterile preferred)
- Previous experience in HAs inspection support (backroom / SME)
- Experienced in QMS document management
- Fluency in English
- Experience working with electronic quality systems (e.g. change controls, deviations, OOX, complaints, etc.)
- Strong quality mindset, documentation, communication, and cross-functional collaboration skills.

Languages:

- Italian

- English

Commitment to Diversity and Inclusion: Novartis is committed to building an outstanding, inclusive work environment and diverse teams' representative of the patients and communities we serve.

## Role Requirements

**Why Novartis:** Helping people with disease and their families takes more than innovative science. It takes a community of smart, passionate people like you. Collaborating, supporting and inspiring each other. Combining to achieve breakthroughs that change patients' lives. Ready to create a brighter future together? <https://www.novartis.com/about/strategy/people-and-culture>

**Benefits and Rewards:** Learn about all the ways we'll help you thrive personally and professionally.

[Read our handbook \(PDF 30 MB\)](#)

Division

DIV\_GD

Business Unit

Quality

Location

LOC\_IT

Site

Ivrea

Company / Legal Entity

IT58 (FCRS = IT058) Advanced Accelerator Applications Italy Srl

Functional Area

FCT\_QA

Job Type

Full time

Employment Type

Regular

Shift Work

No

[Apply to Job](#)

Job ID

REQ-10074537

## TRD Sr. QA Specialist

[Apply to Job](#)

---

**Source URL:** <https://prod1.jobapi.novartis.com.cn/req-10074537-trd-sr-qa-specialist>

### List of links present in page

1. <https://prod1.jobapi.novartis.com.cn/req-10074537-trd-sr-qa-specialist>
2. <https://www.novartis.com/about/strategy/people-and-culture>
3. [https://www.novartis.com/sites/novartis\\_com/files/novartis-life-handbook.pdf](https://www.novartis.com/sites/novartis_com/files/novartis-life-handbook.pdf)
4. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Ivrea/TRD-Sr-QA-Specialist\\_REQ-10074537](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Ivrea/TRD-Sr-QA-Specialist_REQ-10074537)
5. [https://novartis.wd3.myworkdayjobs.com/en-US/Novartis\\_Careers/job/Ivrea/TRD-Sr-QA-Specialist\\_REQ-10074537](https://novartis.wd3.myworkdayjobs.com/en-US/Novartis_Careers/job/Ivrea/TRD-Sr-QA-Specialist_REQ-10074537)